A 219-mer CHO-Expressing Receptor-Binding Domain of SARS-CoV S Protein Induces Potent Immune Responses and Protective Immunity

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Abstract

Development of vaccines is essential for the prevention of future recurrences of severe acute respiratory syndrome (SARS), caused by the SARS coronavirus (SARS-CoV). The spike (S) protein, especially receptor-binding domain (RBD) of SARS-CoV, plays important roles in the prevention of SARS infection, and is thus an important component in SARS vaccine development. In this study, we expressed a 219-mer (residues 318–536) RBD protein in Chinese hamster ovary (CHO)-K1 cells (RBD219-CHO), and tested its immune responses and protective immunity in a mouse model. The results showed that this recombinant protein was correctly folded, being able to maintain intact conformation and authentic antigenicity. It could induce strong humoral and cellular immune responses and high titers of neutralizing antibodies in the vaccinated mice. RBD219-CHO protein elicited potent protective immunity that protected all vaccinated mice from SARS-CoV challenge. These results suggest that the recombinant RBD219-CHO protein has great potential for the development of an effective and safe SARS subunit vaccine.

Introduction

A newly emerging infectious disease, severe acute respiratory syndrome (SARS), is caused by SARS coronavirus (SARS-CoV) (27,46,58), a zoonotic virus that most likely originated in its natural reservoir bats, through intermediate transmission such as via palm civets and raccoon dogs, and was finally transmitted to humans (21,37,38). Transmission of SARS-CoV from humans to humans led to the global outbreak of SARS in 2003 (39,44,45,52). Though SARS is currently under control, it is necessary to develop effective and safe vaccines for the prevention of future SARS outbreaks that may arise from animal reservoirs or accidentally due to laboratory virus escape.

Recently developed SARS vaccines are of various categories (18), including inactivated virus vaccines (49,51,59), subunit vaccines (2,26), DNA vaccines (28,42,55), virus-like particles (40,41), viral vector-based vaccines (4,19,34), and different vaccine combinations (16,30,53). A variety of SARS vaccines have been tested in animals, including monkeys, ferrets, mice, and hamsters (1,7,9,20,33,48,49), and some of them have been evaluated in humans (42). These vaccines may target different antigens of the virus, but most of them are based on the spike (S) protein. It has been reported that an adenovirus-based vaccine expressing S protein prevented pneumonia in ferrets after SARS-CoV challenge, and stimulated potent immune responses in macaques (36). A recombinant SARS S-protein elicits neutralizing antibodies and protection in mice (29). Specific humoral and cellular immune responses and/or protection could be induced by SARS S DNA vaccines via different vaccination routes (28,55). A SARS-CoV-like particle carrying the S protein protected mice from virus challenge (40). These reports suggest that the S protein plays an important role in the prevention of SARS infection (3).

Our previous studies demonstrated that a recombinant fusion protein consisting of a 193-mer (residues 318–510) receptor-binding domain (RBD) of SARS-CoV S protein tagged with the Fc fragment of human IgG (RBD193-Fc) could induce highly potent neutralizing antibody responses and protective immunity (14,25). However, one potential disadvantage of this vaccine candidate is that the Fc tag, which was added to the C-terminus of RBD193 in the hope of increasing immunogenicity by binding Fc-tagged immunogen
Materials and Methods

Gene construction, protein expression, and purification of RBD219-CHO

The gene construction and expression of RBD219-CHO protein was done as previously described (15). The genes encoding the fragment containing 219 aa (318–536) of the SARS-CoV S protein RBD region, plus a 6-His tag at the C terminus, were amplified by PCR using a full-length S plasmid (Tor2 strain) as the template (12, 24). It was then inserted into the GS Gene Expression Vector PEE14.1. The constructed recombinant plasmid (RBD219) was confirmed by sequencing analysis. Briefly, the recombinant RBD219 plasmid was transiently transfected using FuGENE 6 transfection reagents (Roche Applied Science, Indianapolis, IN) into CHO-K1 cells precultured in F-12K medium (American Type Culture Collection, Manassas, VA). The culture medium was replaced by fresh OPTI-MEM I Reduced-Serum Medium (Invitrogen, Carlsbad, CA) 10 h later, and the supernatant was collected 72 h post-transfection. Culture supernatant containing expressed protein was added to protease inhibitor cocktails (Roche Applied Science) and purified using His columns (Promega, Madison, WI). The protein yield using this transient transfection method was about 10.6 mg/g.

Detection of RBD219-CHO protein reactivity with RBD-specific mAbs

The reactivity of purified RBD219-CHO protein was determined by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), followed by Western blot, according to our previously described protocols (10). Briefly, purified protein (2 µg/well) was separated by non-reducing SDS-PAGE (no boiling and no reducing agent). The gel was transferred to nitrocellulose membranes (Bio-Rad Laboratories, Hercules, CA), which were then blocked overnight at 4°C with blocking buffer containing 5% non-fat milk dissolved in 0.1% Tween 20-PBS (PBST). The blots were incubated for 1 h at room temperature with serial RBD-specific monoclonal antibodies (mAbs) generated in our laboratory (25), at a final concentration of 0.5 µg/mL. After washing three times, the blots were incubated with horseradish peroxidase (HRP)-conjugated goat anti-mouse IgG (1:5000; Zymed, Carlsbad, CA) for 1 h at room temperature. Signals were visualized with ECL Western blot Substrate Reagents and Amersham Hyperfilm (GE Healthcare, Piscataway, NJ).

Mouse vaccination and sample collection

A group of five female BALB/c mice 4–6 wk of age were subcutaneously vaccinated with purified RBD219-CHO protein (20 µg/mouse) in the presence of Freund’s complete adjuvant (FCA; Sigma, St. Louis, MO), and boosted twice with the same immunogen (10 µg/mouse) containing Freund’s incomplete adjuvant (FIA; Sigma) at 21-day intervals. Mice injected with the same amount of PBS were used as negative controls. Sera were collected pre-immunization and 10 d after each vaccination, and analyzed for IgG and neutralizing antibody responses. Spleen cells collected 10 d after the last vaccination were used for the detection of cellular responses. For parallel experiments, mice were challenged with SARS-CoV 10 days after the last vaccination, and analyzed for viral load and RNA copies in challenged lung tissues 5 d post-challenge. Vaccination protocols are described in detail in Fig. 1.

Enzyme-linked immunosorbent assay (ELISA)

Mouse sera were analyzed using ELISA for IgG antibody specific for RBD of SARS-CoV S protein using a previously described protocol with some modifications (14). Briefly, 96-well plates were pre-coated with 100 µL/well of RBD219-CHO protein (1 µg/mL) and sat overnight at 4°C, then blocked with 2% non-fat milk at 37°C for 2 h. Serially diluted mouse sera were added to the plates at 100 µL/well and incubated at 37°C for 1 h, followed by three washes with PBST. Bound antibodies were then reacted with HRP-conjugated goat anti-mouse IgG (1:2000) at 37°C for 1 h. The substrate 3,3',5,5'-tetramethylbenzidine (TMB; Zymed) was added to the plates for 8 C and allowed to develop for 30 min. Absorbance was measured at 450 nm.

FIG. 1. Vaccination protocol for the experiments.
Neutralization assay for SARS pseudovirus infection

The neutralization assay against SARS-CoV pseudovirus infection was done according to our previously established method (25). Pseudotyped virus bearing SARS-CoV S protein, and defective HIV-1 genome expressing luciferase as reporter was prepared as before. Briefly, 293T cells were co-transfected with a plasmid encoding codon-optimized SARS-CoV S protein, and a plasmid encoding Env-defective, luciferase-expressing HIV-1 genome (pNL4-3.luc.RE), using FuGENE 6 transfection reagents (Roche Applied Science). Supernatants containing SARS pseudovirus were harvested 72 h post-transfection and used for single-cycle infection of 293T cells expressing ACE2 (ACE2/293T). The cells were plated at 10^5 cells/well in 96-well tissue culture plates 24 h pre-infection. The pseudovirus-containing supernatants were pre-incubated with serially diluted mouse sera at 37 °C for 1 h before they were added to the cells. After 24 h, the cells were re-fed with fresh medium, which was followed by lysing the cells 72 h later using cell lysis buffer (Promega), and the lysates were transferred into 96-well luminometer plates. Luciferase substrate (Promega) was added to the plates, and relative luciferase activity was determined with an Ultra 384 luminometer (Tecan Systems). The neutralization of SARS pseudovirus was calculated (8), and presented as 50% neutralizing antibody titer (NT50).

Neutralization assay for live SARS-CoV infection

Neutralization assay against the live SARS-CoV (GZ50 strain; GenBank accession no. AY304495) was done as previously described (16,17), and neutralizing antibody titers of sera from mice immunized with RBD219-CHO protein were detected in Vero E6 cells. Briefly, the cells were seeded (10^4 cells/well) on 96-well tissue culture plates 24 h before infection. Serial twofold dilutions of serum samples were separately added to the cells in triplicate. Cytopathic effect (CPE) was detected in Vero E6 cells. Briefly, the cells were seeded (10^4 cells/well in 96-well tissue culture plates 24 h before infection. After 24 h, the cells were re-fed with fresh medium, which was followed by lysing the cells 72 h later using cell lysis buffer (Promega), and the lysates were transferred into 96-well luminometer plates. Luciferase substrate (Promega) was added to the plates, and relative luciferase activity was determined with an Ultra 384 luminometer (Tecan Systems). The neutralization of SARS pseudovirus was calculated (8), and presented as 50% neutralizing antibody titer (NT50).

ELISPOT assay

ELISPOT assay was done using a mouse kit (Mabtech Inc., Mariemont, OH), according to the manufacturer’s protocol and our previously described method (16,17). Briefly, 96-well ELISPOT plates were coated with anti-IFN-γ, anti-IL-2, and anti-IL-4 mAbs overnight at 4 °C, and blocked with RPMI-1640 containing 10% FBS for 2 h at room temperature. Single-cell suspensions (2×10^5 cells/well) isolated from vaccinated mouse spleens were added to the wells, which was followed by incubation at 37 °C for 24 h in the presence of an MHC-H-2d-restricted SARS-CoV RBD-specific cytotoxic T lymphocyte (CTL) peptide (N50: S365–374, KCYGVSATKL), or Th peptide (N60: S435–444, NYNYKYRRLY), at a final concentration of 1 μg/mL (30). The plates were washed with PBS, followed by incubation with biotinylated labeled antimouse IFN-γ, IL-2, and IL-4 mAbs at 1:1000 for 2 h at room temperature. After additional washes, the wells were incubated with streptavidin-conjugated HRP for 1 h at room temperature, and developed with TMB substrate. Spots of cytokine-producing T cells were counted by using an ELISPOT reader and ImmunoSpot 3 software (Cellular Technology Ltd., Cleveland, OH). Results were expressed as the number of spot-forming cells per 10^6 input cells.

Challenge of vaccinated mice with live SARS-CoV

Vaccinated mice were intranasally inoculated with SARS-CoV strain GZ50 (5×10^3 TCID50) as previously described (16,17). The mice were sacrificed 5 d after virus challenge, and the lungs were removed and stored at −80 °C for further virological detection.

Quantitative reverse transcriptase polymerase chain reaction (qRT-PCR)

The viral RNA copies in the lung tissues were determined by qRT-PCR according to a previously described protocol (14). Briefly, total RNA was extracted from 20 mg of the lung tissues using a RNeasy Mini kit (Qiagen, Valencia, CA) according to the manufacturer’s instructions. SARS-CoV RNA was quantified in a 30-μL mixture containing 10 μL RNA, 15 μL 2×TaqMan® One-Step RT-PCR Master Mix (Applied Biosystems, Foster City, CA), 0.75 μL 40× multiscribe, 0.25 μM each forward primer A (Taq-772F 5'-AAG CCT CGC CAA AAA CGT AC-3'), reverse primer R (Taq-1000R 5'-AAG CCT CGC CAA AAA CGT AC-3'), reverse primer Ar (Taq-1000R 5'-AAG CCT CGC CAA AAA CGT AC-3'), reverse primer Ar (Taq-1000R 5'-AAG CCT CGC CAA AAA CGT AC-3'), probe (Taq-955T 5'-FAM-TCA GGC ATG GAA -3'), and probe (Taq-955T 5'-FAM-TCA GGC ATG GAA GTC ACA CT-TAMRA) (TIB Molbiol, Berlin, Germany), using a fluorometric PCR instrument (ABI 7300; Applied Biosystems).

Viral titer and total virus detection

The virus replication was detected by titration of the inoculated virus in lung tissues collected from sacrificed mice according to a previously described protocol (14). Briefly, the lung tissues were homogenized to a final concentration of 10% (w/v) suspension in DMEM. Tissue homogenates were centrifuged, filtered, and inoculated into the monolayer of Vero E6 cells seeded on 96-well tissue-culture plates. The results were evaluated after 3 d of culture under phase-contrast microscopy, and viral titers using a CPE-based TCID50 test were calculated by the Reed-Muench method. Viral titers were expressed as log_{10}TCID50/g of tissue, with a lowest detection limit of 1.5 log_{10}TCID50/g. The total amount of virus was calculated by multiplying the weight of the lung tissues and the viral titers measured in 10% of tissue homogenates.

Results

The expressed RBD219-CHO protein was correctly folded and maintained intact conformation, and was recognized by a majority of conformational epitope-specific mAbs against RBD

The expressed RBD protein was analyzed for conformation by using mAbs targeting the conformational epitopes in RBD. As shown in Fig. 2, the expressed protein reacted with
5 of 6 selected mAbs recognizing conformational epitopes, having the strongest reaction with the conformational V mAb 33G4. It also reacted with mAb 17H9, that recognizes the linear epitope in RBD. These results demonstrate that the expressed RBD219-CHO protein is correctly folded and maintains intact conformation.

**RBD219-CHO induced high titers of RBD-specific antibodies in vaccinated mice**

In order to detect humoral immune responses induced by RBD219-CHO protein, sera were collected from vaccinated mice and analyzed with ELISA for RBD-specific antibodies. As shown in Fig. 3A, all sera from 10 d after the first, second, and third vaccinations showed strong antibody responses, with those from the last vaccination demonstrating the highest reactivity. The sera from pre-immunization (Pre), and those from PBS controls had only background levels of antibody responses. To determine the titer of RBD-specific antibodies induced by RBD219-CHO protein, sera collected 10 d after the last vaccination were titrated at a series of fourfold dilutions and A450 was measured. Fig. 3B indicates that RBD219-CHO elicited high levels of RBD-specific antibodies, with the end-point titer of $1:3.3 \times 10^6$. In contrast, only background levels of antibody were detected in the PBS control group (Fig. 3B).

**RBD219-CHO induced highly potent neutralizing antibodies against in-vitro infection by SARS pseudovirus and live SARS-CoV**

Sera collected 10 d after the last vaccination were tested for neutralizing activity against infection by SARS pseudovirus in ACE2/293T cells, and live SARS-CoV in Vero E6 cells. As shown in Fig. 4, $1.58 \times 10^4 \pm 4.9 \times 10^3$ of NT$_{50}$ were reached in RBD219-CHO-vaccinated mouse sera to neutralize SARS pseudovirus infection, levels higher than those of PBS controls. High titers of RBD-specific IgG antibodies were also confirmed to neutralize live SARS-CoV in infected Vero E6 cells, with $1:1.0 \times 10^5 \pm 2.4 \times 10^2$ NT$_{50}$ of neutralizing antibodies maintained in the vaccinated mice (Fig. 4). These results suggest that highly potent neutralizing antibodies may be induced by RBD219-CHO protein against both SARS pseudovirus and live SARS-CoV infection in cell cultures in vitro.

**RBD219-CHO was able to induce a relatively high level of CTL and Th1-/Th-2 responses in vaccinated mice**

For detection of the cellular immune responses potentially induced by RBD219-CHO protein, mouse splenocytes were collected 10 d after the last vaccination, and T-cell responses were detected by ELISPOT. Splenocytes were stimulated with N50 and N60 peptides specific to CD8$^+$ T (CTL) cells and CD4$^+$ T (Th) cells, respectively, in the RBD of SARS-CoV (30). For further detection of the cellular immune response, purified RBD protein was also used for stimulation with a final concentration of 1 $\mu$g/mL. As shown in Table 1, RBD219-CHO induced RBD-specific CTL responses, with a particularly high frequency of IL-2-producing CD8$^+$ (CTL) cells under the stimulation of the CTL peptide N50. Evaluation of Th-cell responses upon stimulation with the N60 peptide indicated that RBD219-CHO protein may elicit relatively higher levels of Th-1 response, as represented by the production of high numbers of IL-2-presenting T cells, in addition to the induction of Th-2 response, which was indicated by the presence of IL-4-secreting T cells in the spleens of vaccinated mice. The results in Table 1 also show that the cellular immune responses induced by RBD219-CHO protein stimulation were generally higher than those elicited by the CTL peptide N50 or Th peptide N60 stimulation alone, probably because the recombinant protein covering the RBD region contains much more CTL and/or Th epitopes. In contrast, only a background level of cellular response was detected in the splenocytes of mice from the PBS controls (Table 1), or the cells stimulated with the supernatants of CHO culture only (data not shown). These data suggest that the expressed RBD219-CHO protein was able to induce a relatively high level of cellular immune response in the vaccinated mice.

**Complete suppression of SARS-CoV replication in lung tissues of virus-challenged mice, possibly mediated by RBD219-CHO-induced neutralizing antibody responses**

To detect the protective immunity induced by RBD219-CHO protein, vaccinated mice were challenged with SARS-CoV GZ50, and mouse lung tissues were collected 5d post-virus challenge to measure virus replication. The mice did not
develop obvious weight loss or clinical signs of disease in both the control and vaccinated groups. No detectable replication was seen in all five mice vaccinated with RBD219-CHO protein. In contrast, high levels of virus replication were found in the lung tissues of PBS control mice (Fig. 5A). The above results confirm that RBD219-CHO protein was able to completely protect vaccinated mice from subsequent SARS-CoV challenge.

To investigate the relationship between neutralizing antibody and virus protection, we compared the serum neutralizing antibody titer and the viral RNA copies in the lung tissues after the mice were challenged with live SARS-CoV (GZ50). As shown in Fig. 5B, no detectable virus replication was accompanied by high titers of neutralizing antibodies (1:1.0×10^3 ± 2.4×10^2) in the RBD219-CHO-vaccinated mice, while high levels of virus replication (1:1.9×10^3 ± 4.1×10^2 RNA copies/µg of lung tissue) and undetectable levels of serum neutralizing antibodies (<1:20) were found in the PBS control mice. These results suggest that neutralizing antibody responses may play a critical role in the inhibition of SARS-CoV replication.

**Discussion**

Vaccines based on the recombinant S protein of SARS-CoV have been shown to induce humoral immune responses, especially neutralizing antibodies against SARS-CoV (23,33,35,60). High titers of cross-neutralizing antibodies against

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**FIG. 3.** Detection of RBD-specific antibodies in sera of mice immunized with RBD219-CHO protein and PBS controls. The data are presented as mean A450 ± standard error (SE) from five mice per group. (a) IgG antibody detected in sera (at 1:3000 dilution) collected before immunization (Pre) and 10 d after each vaccination are shown here. (b) RBD-specific antibody titer detected in sera collected 10 d after the last vaccination are shown here.

**FIG. 4.** Detection of neutralizing antibodies in sera of mice vaccinated with RBD219-CHO or PBS controls. Sera collected 10 d after the last immunization were tested for neutralizing activity against infection by SARS pseudovirus in ACE2/293T cells, and live SARS-CoV in Vero E6 cells. The data are presented as mean NT_{50} ± SE of five mice per group.
pseudovirus in vitro were detected in rRBD-vaccinated mice and rabbits (24). Protective humoral responses were detected in protein-vaccinated rabbits, with antisera against S containing neutralizing antibodies for SARS-CoV infection in vivo (43). These reports demonstrated that S protein-induced neutralizing antibodies play important roles in fighting SARS-CoV infection.

We previously reported that a recombinant protein containing a 193-aa RBD with Fc tag (RBD193-Fc) induced highly potent neutralizing antibody responses and protective immunity in vaccinated animals (14). In order to eliminate any potential adverse effects that might be induced by RBD193-Fc when used as a vaccine component in humans, we have expressed several RBD proteins with the Fc tag removed in mammalian cells 293T (RBD-293T) and CHO-K1 (RBD193-CHO), insect cells Sf9 (RBD-Sf9), and E. coli (RBD-Ec). Unlike RBD proteins expressed in 293T, Sf9, and E. coli, that may induce complete protection against live SARS-CoV challenge in vaccinated mice, the 193-aa RBD protein expressed in CHO-K1 cells (RBD193-CHO) protected a majority of the mice from virus challenge, with virus replication detected in two of five mice (13,15). When we expressed a 219-mer RBD protein in the same cell line of CHO-K1 (RBD219-CHO), the newly-designed protein elicited high titers of RBD-specific antibodies with potent neutralizing activity against both SARS pseudovirus and live SARS-CoV infection in cell cultures in vitro. Like rRBD proteins expressed in 293T, Sf9, and E. coli, RBD219-CHO was able to protect all vaccinated mice against SARS-CoV challenge in vivo (Fig. 5). Similarly to other rRBD proteins expressed above, this mammalian cell CHO-K1-expressed RBD protein without the Fc fusion tag reacted with the majority of the mAbs specific for the conformational and linear epitopes in RBD, suggesting that it maintains intact conformation and proper antigenicity (Fig. 2). Similarly to RBD193-CHO protein, a stable cell line could be established in the CHO-K1 cell lines for long-term expression of the RBD219-CHO protein (data not shown). The RBD219-CHO protein (residues 318–536) was expressed with an extension of 26 residues at the C terminus of the RBD193-CHO protein (residues 318–510). This extended fragment may assist RBD to refold into a more stable or more immunogenic conformation to induce more potent protective immunity in the vaccinated animals. Future comparison of the structure and immune responses and long-term protection of these two proteins is warranted.

In addition to humoral immune responses, cellular immune responses may play a role in the clearance of SARS-CoV infection, in which both CD4+ and CD8+ T cells are involved in the immune response against virus infection. Memory T-cell responses against S protein were shown to continue for more than 1 y after SARS-CoV infection, as indicated by the production of IFN-γ (54). An increased specific

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*Typical ELISPOT detection of cellular immune responses in CHO-K1-vaccinated mice.

Table 1. ELISPOT Detection of Cellular Immune Responses in RBD219-CHO-Vaccinated Mice

![FIG. 5. Detection of protective immunity in mice immunized with RBD219-CHO protein or PBS controls. Mice vaccinated with RBD219-CHO protein were challenged with live SARS-CoV (GZ50). (a) SARS-CoV replication in the lung tissues of challenged mice was detected and expressed as log_{10} TCID_{50}/g of tissue. The detection limit was 1.5 log_{10} TCID_{50}/g of lung tissue. M1–M5 indicate the five mice used per group. (b) The number of SARS-CoV RNA copies in the lung tissues of mice challenged with live SARS-CoV GZ50 were measured by qRT-PCR, and expressed as RNA copies/µg of lung tissue. The neutralizing antibody titers (NT_{50}) in the sera of the mice are also shown. The data are presented as mean ± SE of five mice per group. The experiment was performed three times and similar results were obtained.*
CTL response was induced by an S2 fragment encoding residues 681–980 (22). Immunization of mice with a DNA fragment encoding the S protein was demonstrated to induce both CD4+ and CD8+ T-cell responses (31). In a Phase I human study, an S protein-expressing DNA vaccine elicited SARS-CoV-specific CD4+ T-cell responses in all tested healthy adults, and CD8+ T-cell responses in 20% of individuals (42). Both CTL and Th-1/Th-2 responses were detected in this study in RBD219-CHO-vaccinated mouse splenocytes, more evidence of the high immunogenicity and antigenicity of the expressed RBD protein when used as a vaccine candidate (Table 1).

Although both humoral and cellular immune responses are important for suppression of SARS-CoV infection, neutralizing antibodies may be the key factor for protecting mice from subsequent virus challenge (13,14,50). Long-term protection mediated by neutralizing antibodies could be achieved by immunization of mice with a single dose of an attenuated vesicular stomatitis virus (VSV) recombinant bearing S protein (VSV-S) (34). A recombinant secreted protein containing 14–762 residues of the S protein could provide sufficient neutralizing antibodies to protect against SARS-CoV infection (2). The protective immunity seen in the mice vaccinated with an S protein-expressing DNA vaccine was mediated by humoral immunity rather than by a T-cell-dependent immune mechanism (55). In agreement with the above reports, our results show that neutralizing antibodies induced by RBD219-CHO protein play a significant role in the suppression of SARS-CoV infection (Fig. 5). We and others have demonstrated that RBD of SARS-CoV S protein is the basis for the development of vaccines against SARS-CoV (6,11,16,17,26,32). High titers of neutralizing antibodies and protective immunity could be induced by subunit vaccines based on the RBD protein with or without fusion segments (14,56). However, RBD-based vaccines without the fusion tag would eliminate any potential adverse effects induced by fusion proteins when used in humans (13). In agreement with our previous reports, the results of this study further confirm that the recombinant RBD protein without the Fc fusion fragment is able to induce potent neutralizing antibody responses and protection against infection by SARS-CoV, suggesting the possibility of the development of RBD subunit-based vaccines for SARS prevention.

Acknowledgments

This study was supported by the National Institutes of Health (NIH) of the United States (ROI AI68002), by the Research Fund for the Control of Infectious Diseases, the Health, Welfare and Food Bureau of the Hong Kong SAR Government, and by the National 973 Basic Research Program of China (2005CB523001).

Author Disclosure Statement

No competing financial interests exist.

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Received October 9, 2009; accepted December 17, 2009.